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Claims:

- 1. A method of using ionic liquids such as molten salts as solvents in headspace gas chromatography wherein said method comprises dissolving or dispersing a sample in at least one ionic liquid and volatilizing the volatile components of the sample.
- 5 2. The method according claim 3 wherein the ionic liquid has a melting point of less than 100°C.
 - 3. The method according to claim 2 wherein the ionic liquid has a melting point of less than 30°C.
- The method according to any preceding claim wherein the ionic liquid has a vapor
 pressure of less than about 1 mm/Hg at 25°C.
 - 5. The method according to claim 4 wherein the ionic liquid has a vapor pressure of less than about 0.1 mm/Hg at 25°C.
 - 6. The method according to claim 5 wherein the ionic liquid has essentially no vapor pressure.
- 15 7. The method according to any preceding claim wherein the thermal stability of the ionic liquid is form 150°C to 400° C.
 - 8. The method according to claim 7 wherein the thermal stability of the ionic liquid is from 200° C to 300° C.
- 9. The method according to claim 1 wherein the ionic liquid has a melting point of less than 250°C, a vapor pressure less than about 1mm/Hg at 25°C and the thermal stability of the ionic liquid is from 150° C to 400° C.
 - 10. The method according to any preceding claim wherein the anion of the ionic liquid is selected from the group consisting of Cl⁻, Br⁻, NO₂⁻, NO₃⁻, AlCl₄⁻, BF₄⁻, PF₆⁻, CF₃COO⁻, CF₃SO₃⁻, (CF₃SO₂)₂N⁻, OAc⁻, CuCl₃⁻, GaBr₄⁻, GaCl₄⁻, and SbF₆⁻.
- 25 11. The method according to any preceding claim wherein the cation of the ionic liquid is selected from the group consisting of pyridinium, ammonium, imidazolium, phosphonium, and sulphonium.



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- 12. The method according to any preceding claim wherein the ionic liquid is selected from the group consisting of an imidazolium salt, pyridinium salt, ammonium salt, phosphonium salt, and sulphonium salt, and mixtures thereof.
- 13. The method according to claim 12 wherein the imidazolium salt has formula (I)

$$\begin{array}{c}
 & R^1 \\
 & R^2
\end{array}$$
(I)

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wherein R^1 and R^2 are independently selected from the group consisting of a C_1 - C_{18} aliphatic group and a C_4 - C_{18} aromatic group; and A^- is an anion.

14. The method according to claim 12 wherein the ammonium salt has formula (II)

$$\begin{array}{ccc}
R^{5} \stackrel{R^{6}}{\swarrow} & R^{3} \\
\downarrow & A^{-} \\
R^{4} & A^{-}
\end{array}$$
(II)

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wherein R³, R⁴, R⁵ and R⁶ are independently selected from the group consisting of a C₁-C₁₈ aliphatic group and a C₄-C₁₈ aromatic group; and A is an anion.

15. The method according to claim 12 wherein the phosphonium salt has formula (III)

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$$\begin{array}{ccc}
R^{10} & & \\
R^{5} & & \\
R^{8} & & A^{-}
\end{array}$$
(III)

wherein R⁷, R⁸, R⁹, and R¹⁰ are independently selected from the group consisting of a C₁
20 C₁₈ allphatic group and a C₄-C₁₈ aromatic group; and A⁻ is an anion.

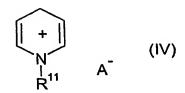
16. The method according to claim 12 wherein the pyridinium salt has formula (IV)

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wherein R11 is selected from the group consisting of a C1-C18 aliphatic group and a C4-C18 aromatic group; and A is an anion.

- The method according to any preceding claim wherein the ionic liquid is selected 5 17. from the group consisting of 1-butyl-3-methylimidazolium hexafluorophosphate, 1-hexyl-3methylimidazolium hexafluorophosphate, 1-octyl-3-methylimidazolium hexafluorophosphate, 1-decyl-3-methylimidazolium hexafluorophosphate, 1-dodecyl-3-methylimidazolium hexafluorophosphate, 1-ethyl-3-methylimidazolium bis((trifluoromethyl)sulphonyl)amide, 1hexyl-3-methylimidazolium bis((trifluoromethyl)sulphonyl)amide, 1-hexylpyridinium 10 tetrafluoroborate, 1-octylpyridinium tetrafluoroborate, 1-butyl-3-methylimidazolium tetrafluoroborate, 1-methy-3-ethyl imidazolium chloride, 1-ethyl-3-butyl imidazolium chloride, 1-methy-3-butyl imidazolium chloride, 1-methy-3-butyl imidazolium bromide, 1-methyl-3propyl imidazolium chloride, 1-methy-3-hexyl imidazolium chloride, 1-methy-3-octyl imidazolium chloride, 1-methy-3-decyl imidazolium chloride, 1-methy-3-dodecyl imidazolium 15
 - chloride, 1-methy-3-hexadecyl imidazolium chloride, 1-methy-3-octadecyl imidazolium... chloride, 1-methy-3-octadecyl imidazolium chloride, ethyl pyridinium bromide, ethyl pyridinium chloride, ethylene pyridinium dibromide, ethylene pyridinium dichloride, butyl pyridinium chloride, benzyl pyridinium bromide, and mixtures thereof.
- The method according to claim 17 wherein the ionic liquid is selected from the group 20 18. consisting of 1-octyl-3-methyl-imidazolium hexafluorophosphate, 1-hexyl-3-methyimidazolium hexafluorophosphate, 1-butyl-3-methyl-imidazolium hexafluorophosphate, 1butyl-3-methyl-imidazolium tetrafluoroborate, 1-butyl-3-methyl-imidazolium trifluoromethanesulfonate, 1-ethyl-3-methyl-imidazolium trifluoromethanesulfonate, and 1ethyl-3-methyl-imidazolium bis-(trifluoromethanesulfonyl)-amide. 25
 - The method according to any preceding claim to detect volatile components in a 19. sample by headspace gas chromatography, wherein said method comprises dissolving or

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dispersing a sample in at least one ionic liquid and volatilizing the volatile components of the sample.

- 20. The method according to any preceding claim to identify volatile components in a sample by headspace gas chromatography, wherein said method comprises dissolving or dispersing a sample in at least one ionic liquid and volatilizing the volatile components of the sample.
- 21. The method according to any preceding claim to quantify volatile components in a sample by headspace gas chromatography, wherein said method comprises dissolving or dispersing a sample in at least one ionic liquid and volatilizing the volatile components of the sample.
- 22. The method according to any preceding claim wherein the sample is a pharmaceutical compound.
- The method according to any preceding claim to detect impurities in a pharmaceutical compound by headspace gas chromatography, wherein said method
 comprises dissolving or dispersing a pharmaceutical compound in at least one ionic liquid and volatilizing the volatile components of the pharmaceutical compound.

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